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**AMENDMENTS TO THE CLAIMS**

1. (Currently amended) ~~Use of a~~A method of treating a disorder in which aberrant cell division occurs in a human or animal comprising administering to said human or animal a therapeutically effective amount of a peptide comprising the amino acid sequence

$X_1 X_2 X_3 W M X_4 X_5 X_6 X_7$

wherein

the sequence  $X_1$  to  $X_7$  is an amino acid sequence comprising at least 9 amino acids, which may optionally be interrupted by one or two amino acid residues between one or more of the 9 amino acid positions defined herein;

$X_1$  is selected from W, T, PE, KQI, VV, PQT, H, RI and absent;

$X_2$  is an amino acid with an aromatic side chain;

$X_3$  is P or D;

$X_4$  is an amino acid with a basic side chain;

$X_5$  is an amino acid with a charged side chain;

$X_6$  is an amino acid with a charged side chain; and

$X_7$  is an amino acid with a basic side chain or Serine;

~~in the manufacture of a medicament for treating or preventing a disorder in which aberrant cell division occurs.~~

2. (Currently amended) ~~Use~~The method according to claim 1 wherein  $X_2$  is Y, F or W.

3. (Currently amended) The method ~~Use~~ according to claim 1 or 2 wherein  $X_4$  is K, R or H.

4. (Currently amended) ~~The method~~ Use according to ~~any one of the preceding claims 1~~ wherein X<sub>5</sub> is K, R, E, H, D, N or Q.
5. (Currently amended) ~~Use The method~~ according to ~~any one of the preceding claims 1~~ wherein X<sub>6</sub> is K, R, E, H, D, N or Q.
6. (Currently amended) ~~Use The method~~ according to ~~any one of the preceding claims 1~~ wherein X<sub>7</sub> is H, S, R or K.
7. (Currently amended) ~~Use The method~~ according to claim 1 wherein X<sub>2</sub> is F or Y, X<sub>4</sub> is K or R, X<sub>5</sub> is K, R or E, X<sub>6</sub> is H, R, Q or K and X<sub>7</sub> is H, S, R or K.
8. (Currently amended) ~~Use The method~~ according to claim 7 wherein X<sub>2</sub> is Y and X<sub>3</sub> is P.
9. (Currently amended) ~~Use The method~~ according to claim 8 wherein said peptide X<sub>1</sub> to X<sub>7</sub> has the amino acid sequence W Y P W M K K H H R.
10. (Currently amended) ~~Use The method~~ according to ~~any one of the preceding claims 1~~ wherein said peptide further comprises a cell penetration moiety.
11. (Currently amended) ~~Use The method~~ according to claim 10 wherein said cell penetration moiety is linked directly to the carboxy- terminal of the peptide X<sub>1</sub> to X<sub>7</sub>.
12. (Currently amended) ~~Use The method~~ according to claim 10 or 11 wherein said cell penetration moiety has the amino acid sequence:

X<sub>8</sub> Q I K I W F Q N R R M K W K K

wherein X<sub>8</sub> is R or Q.

- 13 (Currently amended) ~~Use The method~~ according to claim 10 or 11 wherein said cell penetration moiety has the amino acid sequence

X<sub>8</sub> Q X<sub>9</sub> X<sub>10</sub> X<sub>11</sub> W F Q N X<sub>12</sub> X<sub>13</sub> M X<sub>14</sub> W X<sub>15</sub> X<sub>16</sub>

wherein

X<sub>8</sub> is R or Q,

X<sub>9</sub>, X<sub>11</sub> are each independently I or L, and

X<sub>10</sub>, X<sub>12</sub>, X<sub>13</sub>, X<sub>14</sub>, X<sub>15</sub> and X<sub>16</sub> are each independently K or R

14 (Currently amended) ~~Use~~ The method according to claim 10 ~~or 11~~ wherein said cell penetration moiety has the amino acid sequence:

QIRIWFQNRRMKWKK;

QIKIWFQNKRMKWKK;

QIKIWFQNKKMKWKK;

QIRIWFQNRKMKWKK;

QIRIWFQNRRMRWKK;

QIRIWFQNRRMKWRK;

QIRIWFQNRRMKWKR;

QIRIWFQNRRMKWRR;

QIRIWFQNRRMKWKK;

QIKIWFQNRRMKWRK;

QIRIWFQNKRMKWRK;

QIKLWFQNRRMKWKK,

QLKLWFQNRRMKWKK; or

QLRIWFQNRRMKWKK.

15. (Currently amended) ~~Use~~ The method according to claim 10 wherein said peptide has the sequence

W Y P W M K K H H R Q I K I W F Q N R R M K W K, or

W Y P W M K K H H R Q I K I W F Q N R R M K W K K

16. (Currently amended) ~~Use~~ The method according to claim 1 wherein said peptide has the sequence

W Y P W M K K H H R.

17. (Currently amended) ~~Use~~ The method according to ~~any one of the preceding claims~~ wherein said disorder is a cancer.

18. (Currently amended) ~~Use~~ The method according to ~~any one of the preceding claims~~ 1 wherein said cells express one or more Hox genes.

19. (Currently amended) ~~Use~~ The method according to ~~any one of the preceding claims~~ 1 wherein PBX does not act as an oncogene in said cells.

20. (Canceled)

21. (Canceled)

22. (Canceled)

23. (Currently amended) A method reducing the side effects of a cytotoxic or chemotherapeutic agent, in a human or animal comprising administering to said human or animal ~~Use of a~~ the peptide as defined in any one of claims 1 to 16 in the manufacture of a medicament for reducing the side effects of a cytotoxic or chemotherapeutic agent.

24. (Currently amended) A method of maintaing or expanding a stem cell population *in vivo* in a human or animal comprising administering to said human or animal ~~Use of a~~ the

peptide as defined in ~~any one of claims 1 to 16 in the manufacture of a medicament for maintaining or expanding a stem cell population *in vivo*.~~

25. (Canceled)

26. (Currently amended) A method according to claim ~~25~~1 wherein said human or animal is also administered a cytotoxic or chemotherapeutic agent.

27. (Currently amended) A method of maintaining or expanding stem cells *ex vivo* comprising contacting said stem cells with a the peptide as defined in ~~any one of claims 1 to 16.~~

28. (Canceled)

29. (Canceled)

30. (Canceled)

31. (Canceled)

32. (Canceled)

33. (Canceled)

34. (Currently amended) A pharmaceutical composition comprising a peptide as defined in ~~any one of claims 1 to 16~~ and a pharmaceutically acceptable carrier.

35. (Currently amended) A pharmaceutical composition according to claim ~~32~~34 further comprising a cytotoxic or chemotherapeutic agent.